

# Organic geochemistry, stable isotopes, and facies analysis of the Early Aptian OAE—New records from Spain (Western Tethys)

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## ABSTRACT

The Early Aptian Oceanic Anoxic Event (OAE1a) is a time interval characterized by increased organic carbon accumulation in marine sediments, notable sedimentary and biotic changes, and abrupt carbon-isotope excursions indicative of significant major palaeoenvironmental changes linked to a perturbation in the global carbon cycle. Here we present the study of four sections recording the OAE1a (Early Aptian) in Spain, which are located in two broad basins respectively located in the South and the North of Iberia: the Southern Iberian Palaeomargin (Carbonero, La Frontera and Cau sections) and the Basque-Cantabrian Basin (Puentenansa section), which represent depositional settings ranging from shallow marine (distal ramp -Cau- and drowned platform -Puentenansa-) to pelagic environments (Carbonero, La Frontera). Biomarker compositions, C-isotope profiles, biostratigraphic data and facies analysis from the four sections are correlated and integrated. The C-isotope curves all present a clear negative excursion followed by a positive shift. The integration of the C-isotope curves with the biostratigraphic data has been used to correlate the studied sections and to tentatively identify the eight segments formerly proposed from the Alpine domain, and subsequently identified in sections worldwide. Four main groups of compounds are present in all sections: *n*-alkanes, isoprenoids, hopanes and steranes. *n*-Alkanes and acyclic isoprenoids (pristane and phytane) are dominant in most samples. The hopanes are represented by a range of C<sub>27</sub> to C<sub>35</sub> components, with the specific isomers varying amongst the sections due to differences in thermal maturity. Steranes occur as a range of C<sub>27</sub>, C<sub>28</sub> and C<sub>29</sub> isomers, whereas diasteranes only occur in the most thermally mature section (Carbonero). Other compounds of interest include gammacerane and dinosterane. Differences in thermal maturity appear to be the first order control on different biomarker assemblages amongst the studied sections. The Carbonero section is thermally mature, whereas the nearby La Frontera and Cau sections are immature. Puentenansa has intermediate values. Organic matter is derived from a range of terrestrial, marine and bacterial sources. The dominance of the C<sub>29</sub> sterane isomers in all sections suggests a strong contribution from higher plants. The presence of gammacerane indicates water column stratification, and high C<sub>29</sub>/C<sub>30</sub> hopane ratios suggest anoxia at the water/sediment interface, respectively. Sedimentologic analysis also suggests anoxic conditions during sedimentation, but evidence for strong and persistent water column anoxia is equivocal. The correlation of the sections reveals that sedimentation of organic-rich facies started earlier in pelagic and later in the shallow marine settings, which can be related to an expansion of the favorable conditions for organic matter accumulation and preservation from deep marine waters to shallower platform environments during the development of OAE1a.

## Keywords:

Early Aptian  
Oceanic Anoxic Event 1a  
Biomarkers  
C-isotopes  
Spain

## 1. Introduction

The Cretaceous was generally characterized by a greenhouse climate state (Hallam, 1985; Wilson and Norris, 2001), with a reduced latitudinal temperature gradient (Huber et al., 1995; Hay, 2008), elevated pCO<sub>2</sub> levels (Beerling and Royer, 2002, and references therein), and high sea level (e.g. Jenkyns, 1980; Skelton, 2003). In this context, significant short-term changes took place in the ocean-climate

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system, primarily reflected in the widespread deposition of organic carbon-rich marine sediments, whose relation to anoxic marine conditions led to the definition of Oceanic Anoxic Events (Schlanger and Jenkyns, 1976; Jenkyns, 1980). Subsequent research on OAEs have revealed that they represent notable perturbations in the global carbon cycle, which have been related to massive submarine volcanism, formation of oceanic plateaus, and increased rates of seafloor spreading (Leckie et al., 2002). These perturbations were reflected in rapid climate and environmental changes (Jenkyns, 2003; Dumitrescu et al., 2006; Hermoso et al., 2009) affecting both the marine and continental realms (e.g. Skelton, 2003).

Among the OAEs that punctuated the Cretaceous Period, the one that occurred during the Early Aptian (so called OAE1a) was one of the first identified and remains one of the most widely studied. It had a global extent (Arthur et al., 1990; Leckie et al., 2002) and a duration of 1.0 to 1.3 Myr (Li et al., 2008). Sediments recording this event have been recognized in the Tethys domain ("Selli level", Menegatti et al., 1998; Hochuli et al., 1999; Luciani et al., 2001; de Gea et al., 2003; Erba and Tremolada, 2004; Heimhofer et al., 2004; Aguado et al., 2008; de Gea et al., 2008b; Najarro and Rosales, 2008b; Rosales et al., 2009; Mehay et al., 2009; Millán et al., 2009; Najarro et al., 2011a, b; Stein et al., 2011, among others), in the Boreal domain (Gröcke et al., 1999; Föllmi et al., 2006), in the Pacific (Jenkyns, 1995; Ando et al., 2002; Price, 2003; Dumitrescu and Brassell, 2005), and Mexico (Bralower et al., 1999). Dramatic environmental changes occurred in association with OAE1a in settings ranging from continental to pelagic, and particularly in shallow marine carbonate platforms. Among the most striking time-related phenomena were the "nannoconid crisis" (Erba, 1994; Aguado et al., 1999; Erba et al., 2010), profound changes in the marine biota (e.g. Bralower et al., 1994; Erbacher et al., 2001; Leckie et al., 2002), and drastic perturbations in land vegetation and in continental weathering (Föllmi et al., 2006). Nevertheless, other studies suggest that major  $p\text{CO}_2$  and vegetation changes did not occur during OAE1a (Heimhofer et al., 2004).

The impact of OAE1a on shallow carbonate platform environments has been analyzed in several studies. One major environmental perturbation is the widespread drowning event recorded at multiple sites (i.e. Weissert et al., 1998; Föllmi et al., 2006; Castro et al., 2008; Najarro et al., 2011a; Castro et al., 2012). In addition, notable faunal and facies changes in carbonate platforms have been related to the OAE1a (i.e. Huck et al., 2012), including the remarkable widespread development of *Lithocodium*–*Bacinnella* episodes in the Tethys domain (Immenhauser et al., 2005; Bover-Arnal, 2010; Rameil et al., 2010).

A pervasive feature of OAE1a stratigraphic successions is the series of pronounced isotopic excursions, first negative and then positive, that occur in both  $\delta^{13}\text{C}_{\text{carb}}$  and  $\delta^{13}\text{C}_{\text{org}}$  (e.g. Menegatti et al., 1998; Leckie et al., 2002; de Gea et al., 2003; Herrle et al., 2004, among others). The widespread documentation of these features of the carbon isotope profile in numerous successions of marine carbonate and pelagic sediments from the aforementioned Tethys, Boreal and Pacific regions enables their use as a tool for stratigraphic correlation and as a record of the global perturbations of the carbon cycle. Although there is general agreement that the positive excursion in sedimentary  $\delta^{13}\text{C}$  values arises from increased burial of organic matter during the OAE, multiple explanations have been proposed to explain the negative carbon isotope excursion at the onset of OAE1a and, by extension, the causal mechanism. One explanation is that emplacement of large igneous provinces (LIPs; e.g. Ontong-Java and Manihiki Plateaus: Larson, 1991; Larson and Erba, 1999; Tejada et al., 2009; Mehay et al., 2009) raised atmospheric  $p\text{CO}_2$  and enhanced nutrient fluxes to the ocean. Other potential triggers include dissociation of gas hydrates releasing methane in continental margin sediments (Beerling and Royer, 2002; Jahren et al., 2005), or increased recycling rates of  $^{12}\text{C}$  and nutrient-rich intermediate waters linked

to changes in ocean productivity (Menegatti et al., 1998; Larson and Erba, 1999; Erba and Tremolada, 2004; Weissert and Erba, 2004).

This paper presents the study of four sections recording the OAE1a in Spain, from two different basins (Southern Iberian Palaeomargin and North Cantabrian Basin, Fig. 1), and also representing different palaeogeographic settings (shallow marine and pelagic settings). The sections have been characterized using a combination of isotopic, elemental and biomarker approaches. The latter have been widely used in the investigation of OAEs, including the assessment of marine productivity, terrestrial vs. marine sources of organic matter, and the redox state of the ocean (Meyers, 1997; Pancost et al., 2004). The timing of C-rich sediment deposition is interpreted with respect to the local carbon isotope stratigraphy, with biomarker and elemental analyses being used to further constrain changes in organic carbon inputs and differences in thermal maturity among the four sections. Overall, the integration of stratigraphy and geochemistry, and the correlation with reference sections, has led to present a sedimentary model which assess mechanisms for the spread of anoxia, taking into consideration global and regional effects.

## 2. Geological setting

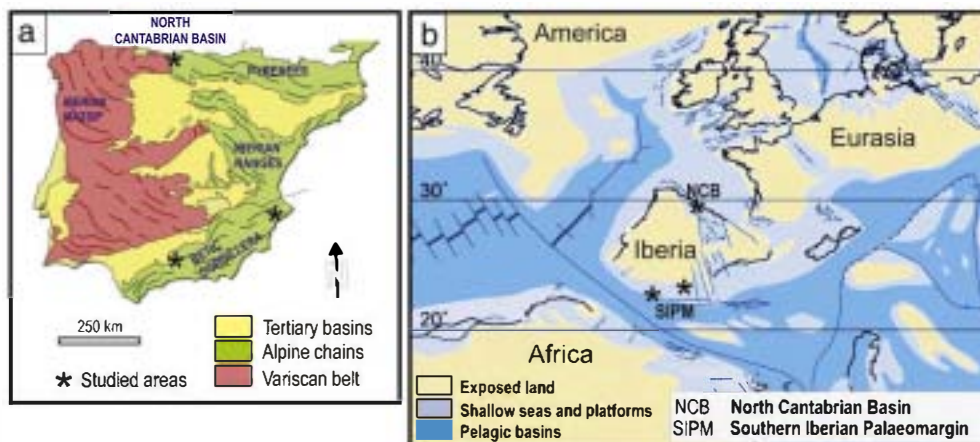
### 2.1. Palaeogeographic framework

During Aptian times the sedimentary basins of Iberia were strongly influenced by the relative motions of the contiguous Eurasian and African plates (Fig. 1). The initiation of the seafloor spreading in the North Atlantic, which started very early in the Cretaceous, led to a decrease in the sinistral movement between Iberia and Africa that had prevailed during part of the Jurassic (Ziegler, 1988), and to a phase of rapid anti-clockwise rotation of Iberia relative to Europe that would culminate in seafloor spreading in the Bay of Biscay from middle Aptian times onwards (Olivet, 1996; Vergès and García-Senz, 2001). In that geodynamic framework, extensional tectonics prevailed in the main sedimentary basins, including the two considered in this paper: the North Cantabrian Basin (NCB), which belonged to the northern margin of the Iberian plate, and the Southern Iberian Palaeomargin (SIPM) (Fig. 1). That extensional tectonism experienced a strong phase during the latest Jurassic to the Hauterivian, followed, during the Barremian–Aptian, by an interval of smaller tectonic movements but larger subsidence rates, that favored the development of wide and thick carbonate platforms (including the Urgonian facies) in the shallow areas of the basins (e.g., Martín-Chivelet et al., 2002; Vera, 2004).

The Lower Cretaceous sequence of the SIPM includes thick successions (> 3000 m) of carbonates and siliciclastics that were deposited in shallow platforms (mostly in the so-called Prebetic Zone, Fig. 2A) and hemipelagic/pelagic settings (dominant in the Subbetic Zone, Fig. 2A) (e.g. Ruiz-Ortiz, 1980; Martín-Chivelet et al., 2002; Vera, 2004). The configuration of the SIPM was defined by a series of basin-scale troughs and swells, bounded by large extensional faults roughly parallel to the continental margin, that were initiated in the Middle Jurassic (e.g., Azéma et al., 1979; García Hernández et al., 1980; Vera, 1988; Ruiz-Ortiz et al., 2001). This tectonic pattern controlled strong differential subsidence and deposition rates. The stratigraphic sections herein considered correspond to depocentral areas in both the Prebetic (Cau section), and the Subbetic (Carbonero and La Frontera sections) (Fig. 2A).

The North Cantabrian Basin (NCB) was a relatively small ( $\approx 20 \times 80$  km), E–W elongated sub-basin, which belonged during the Mesozoic to the larger Basque–Cantabrian Basin (BCB). Located in the northwestern margin of the BCB, this sub-basin behaved independently for most of the Cretaceous, and was relatively less subsident than other areas of the BCB. It was separated from the rest of the basin to the east by a N–S complex fault structure (Río Miera Flexure; Feuillée and Rat, 1971). The NCB was generated by rifting tectonics linked to the opening of the Bay of Biscay and





**Fig 1.** a. Schematic geological map of the Iberian Peninsula showing the main geological units and the location of the studied sections (see Fig. 2 for details). b. Paleogeographic reconstruction of Iberia and its surroundings during the Aptian (simplified from Masse et al., 2000) showing the palaeogeographic location of the studied basins: the Southern Iberian Palaeomargin and the North Cantabrian Basin, respectively located in the southern and the northern margins of Iberia. Note that these basins were inverted and deformed in the Alpine Orogeny during the Tertiary, and are now respectively incorporated into the External Zones of the Betic Cordillera and the Western Pyrenees.

North Atlantic during the Late Jurassic and the Early Cretaceous. This tectonism structured a series of horsts and grabens, mainly outlined by the presence of N-S, NE-SW, and E-W oriented faults, which controlled differential subsidence and sedimentation patterns at least during Barremian-Albian times (Najarro et al., 2007, 2011a). The Lower Cretaceous sediment infill includes relatively thick successions (up to 2000 m thick) of siliciclastics and carbonates, deposited in environments that ranged from fluvial/estuarine to open marine. The studied section corresponds to the Puenteansa road outcrop, near the village of Rábago (Fig. 2B). This area, characterized by moderate subsidence during sedimentation, corresponded to the footwall block of the N-S Bustriguado synsedimentary fault, which is considered as a Cretaceous left-lateral transfer fault (García-Senz and Robador, 2009; Najarro et al., 2009).

Interestingly, the studied successions in the SIPM and the NCB, despite their proximity from a global perspective, were isolated from each other during deposition because of the palaeogeography of Iberia and the palaeoceanography of the surrounding seas. The SIPM belongs to the westernmost part of the northern margin of the Tethys, whereas the NCB belongs to the North Atlantic realm. That relative isolation favored, for example, the development of separate faunal (sub)bioprovinces during Cretaceous times (e.g., Caus and Hottinger, 1986; Caus et al., 2009). Beyond regional tectonics and palaeogeography, the sedimentation in the basins of Iberia should be also influenced in Aptian times by eustatic changes (e.g., Immenhauser et al., 1999; Immenhauser, 2005) and

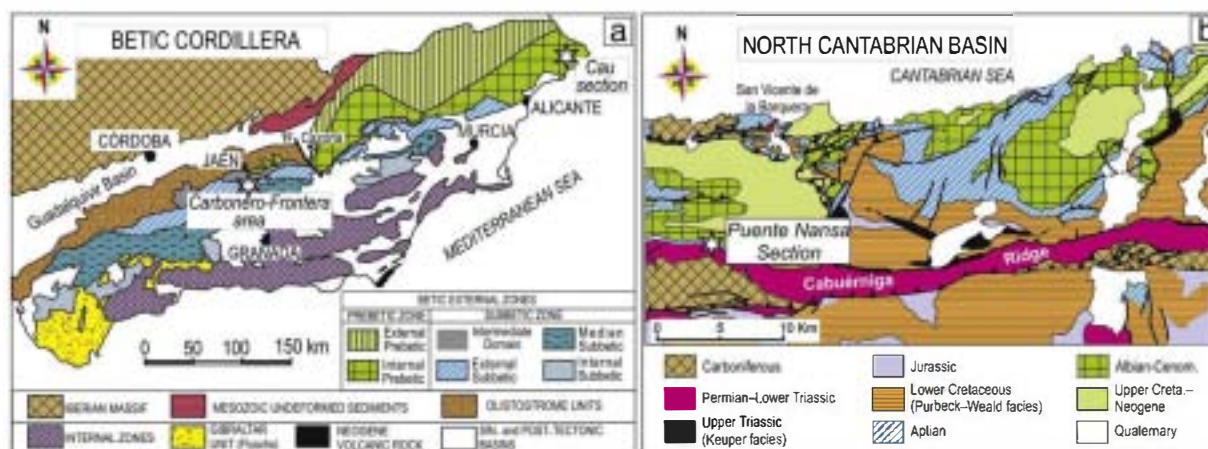
regional climate. At that time Iberia was located approximately between 23° and 33° North latitude, with a tropical to subtropical climate (e.g., Masse et al., 2000).

## 2.2. Previous stratigraphic data of the studied sections

### 2.2.1. Southern Iberian Palaeomargin (SIPM)

Three sections have been previously studied, two from the pelagic Subbetic domain (La Frontera and Carbonero), and one other from the Prebetic domain (Cau section), which represents shallow marine settings.

**2.2.1.1. La Frontera section.** This section is located in the South of the Jaén province (Fig. 2A) and was deposited in a pelagic environment, in a rapidly subsiding area bounded by extensional faults. In this area of the Subbetic Zone, the Barremian is represented by a pelagic rhythmite of the Carretero Formation, whereas the Aptian is represented by the Carbonero Formation, comprising dark marls with an interval of black shales in the Early Aptian, interpreted as the local record of OAE1a (Fig. 3). Here, the presence of calcareous nannofossils has led to the identification of the *Hayesites irregularis* and lower *Rhagodiscus angustus* biozones (Fig. 3). A detailed biostratigraphic characterization of this section, based on planktonic foraminifers, radiolarians and ammonites, can be found in Aguado et al. (1992a) and de Gea (2004).



**Fig. 2.** Location of the studied sections. A.—Betic Cordillera: Cau section belongs to the Prebetic Zone, and Carbonero and La Frontera sections are located in the Subbetic Zone. B.—North Cantabrian Basin: Puenteansa section is located in La Florida area, which formed in the south-western margin of the Basin.

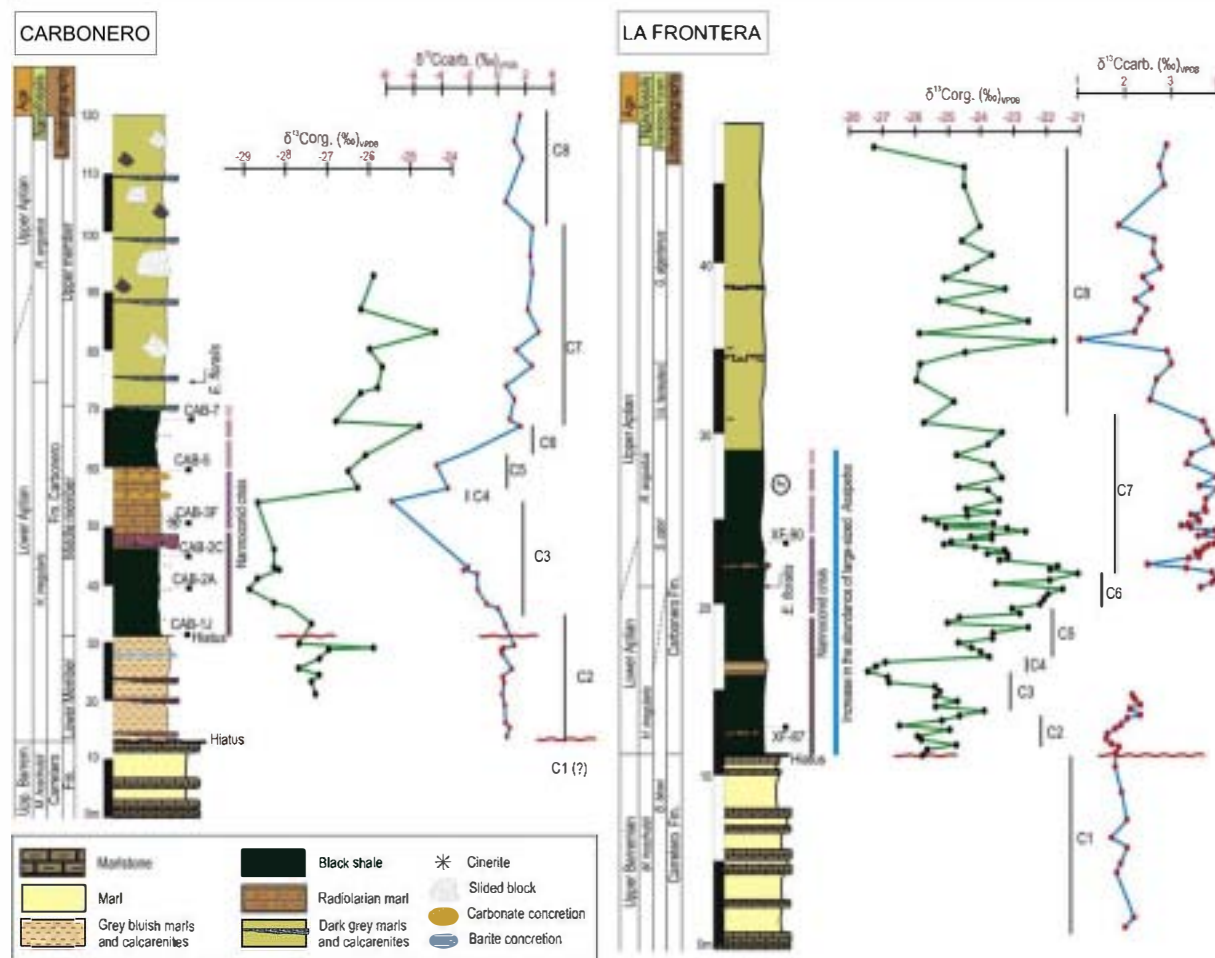
**2.2.1.2. Carbonero section.** This section is located in the South of the Jaén province, fifteen kilometres westward of the La Frontera section. In spite of the proximity, the Carbonero Formation here has strong differences to the La Frontera section and occurs as a rich association of facies, including marls and marlstones, black shales, radiolarian marls, siliceous shales, calcareous turbidites, cinerites, and concretions of barite and carbonate. de Gea et al. (2008a) provided a detailed study of several sections within the area of Carbonero section. They proposed a model of a small subsiding pelagic sub-basin bounded by active faults leading to scarps feeding the turbidite levels, associated with volcanic activity. This small sub-basin has a more complete stratigraphic record than the surrounding areas, with a locally restricted environment leading to high contents of organic matter, along with the development of barite concretions, interpreted to reflect anoxic conditions (Molina and Hernández-Molina, 1993; Bréheret and Brumsack, 2000; de Gea et al., 2008a). Characterization of nannofossil assemblages has led to the recognition of the *Micrantholitus hoschulzi*, *H. irregularis* and lower part of *R. angustus* biozones (Aguado et al., 1992b; Molina et al., 2001; Fig. 3).

**2.2.1.3. Cau section.** This section is located in the NE of the Alicante province (Fig. 2A) and represents the distal parts of a shallow carbonate ramp. In this setting, hemipelagic sedimentation took place during the late Barremian to late Aptian, when a progradational episode led to the deposition of shallow platform carbonates during the latest Aptian–Albian. The late Barremian to late Aptian is represented by the Almadich Formation, comprising hemipelagic marls and marlstones with ammonites, planktonic foraminifers and

calcareous nannofossils. This formation includes a horizon of black marls corresponding to the Early Aptian OAE1a (Fig. 4). This black marl interval occurs in the lower part of the *Schackoina cabri* planktonic foraminifer biozone, *H. irregularis* nannofossil biozone, and corresponds also to the ‘nannoconid crisis’ (Aguado et al., 1999; de Gea et al., 2003). The high organic matter content along with the planktonic foraminifers association (with elongation of the chambers) have been proposed to be related to anoxic–dysoxic conditions in the water column during sedimentation (Aguado et al., 1999; de Gea et al., 2003). The stratigraphy, biostratigraphy and isotope chemostratigraphy of the Cau section have been presented in detail in previous publications (Aguado et al., 1999; de Gea et al., 2003; Castro et al., 2008).

## 2.2.2. North Cantabrian Basin (NCB)

The Puentenansa section is located to the West of Cantabria, in the eastern edge of the North Cantabrian Basin. It is well exposed alongside the Puentenansa road. In this outcrop, the Aptian succession lies unconformably on the Permian to Early Triassic sedimentary rocks. The Early Aptian lithostratigraphy in this area is complex, including four formations: the Rábago and Umbrera Formations (lower Bedoulian), which consist of shallow platform limestones, marls and sandstones; the Párcocinio Formation (mostly early Bedoulian, *Deshayesites weissi* ammonite biozone and middle upper part of the *H. irregularis* nannofossil biozone; Rosales et al., 2009; Najarro et al., 2011b), comprising open marine, black marls and marly siltstones. These deposits, which represent the local record of the OAE1a, reflect the shutdown of the shallow water carbonate factory, as the result of a combined



**Fig. 3.** Carbonero and La Frontera sections. Lithostratigraphy, biostratigraphy, and C-isotope curves. C1 to C8 refers to C-isotope stratigraphy proposed by Menegatti et al., 1998. Samples referred in sections are those selected for biomarker analysis.



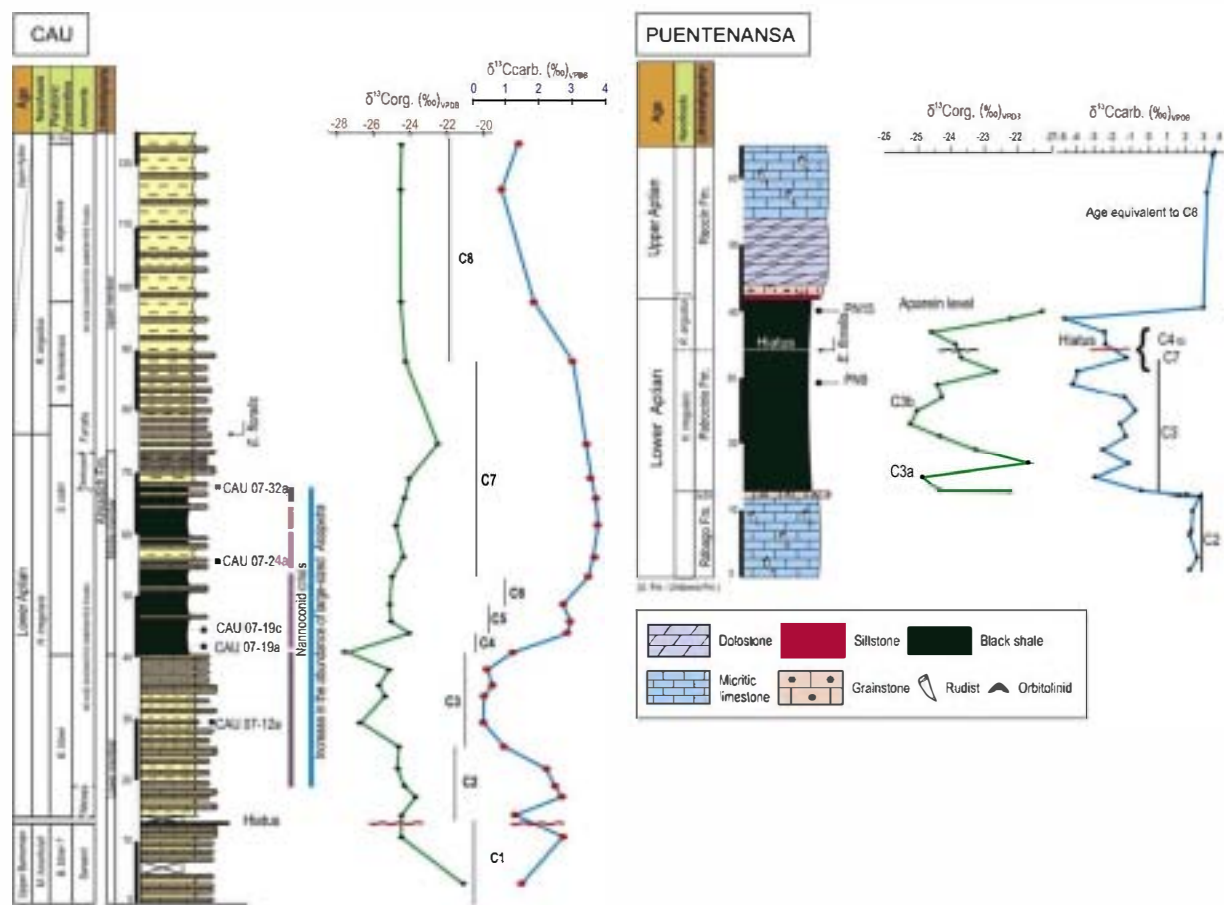


Fig. 4. Cau and Puentenansa sections. Modified from de Gea et al. (2003) and Najarro et al. (2011a). Samples named in sections refer to those selected for biomarker analysis.

action of a relative sea-level rise and environmental stress, and record a net deepening-upward trend leading to a drowned platform (Najarro et al., 2011a). Nannofossil assemblages from the Patrocinio Formation allow the recognition of the *H. irregularis* and lower *R. angustus* biozones (Fig. 4), showing the presence of a stratigraphic gap in the Patrocinio Formation of this area that lasts until the lower Gargasian and which affects the record of OAE1a (Najarro et al., 2011b). Finally, in this section the Reocín Formation (Gargasian), comprising shallow platform limestones, overlies the Patrocinio Formation.

### 3. Methods

#### 3.1. Stratigraphy and biostratigraphy

Through previous extensive field work, the selected sections have been intensively investigated. As a result, the lithology, biostratigraphy and sedimentology of each section has been previously described in detail and finally integrated in a regional stratigraphic context (Aguado et al., 1992a,b, 1999; Castro, 1998; Molina et al., 2001; de Gea et al., 2003; de Gea, 2004; de Gea et al., 2005, 2008b, 2008c; Najarro et al., 2007; Najarro and Rosales, 2008a,b; Najarro et al., 2011a,b).

To allow further and integrated analyses, and to refine the dating and correlation between the different sections, all four were resampled for geochemical analyses and calcareous nannofossil and planktonic foraminiferal characterisation. The marly lithotypes were preferred for sampling in order to obtain the best preserved assemblages. In the Carbonero section, about 33 samples were collected at irregularly spaced intervals (50 cm–5 m) along a 120 m thick succession of marls, marly limestones, black shales and radiolarian marls (Carbonero Formation), spanning the Lower Aptian–lower part of the Upper Aptian interval. The La Frontera section was

sampled at a higher resolution, also at irregularly spaced intervals (15–250 cm), in order to characterize biostratigraphically and sedimentologically the small changes in facies and/or lithology. A total of 100 samples were collected from the marls, marly limestones and black shales of the Carretero and Carbonero Formations spanning the uppermost Barremian–lower part of the Upper Aptian interval. The Cau section was re-sampled for more detailed dating and for geochemical analyses. Finally, in the Puentenansa section, 15 samples of marls and marly siltstones were collected across the Lower Aptian Patrocinio Formation at approximately regular intervals of 2 m, and another 10 limestones were collected from the platform units below and above the Patrocinio Formation.

For calcareous nannofossil characterization, simple smear slides were prepared following standard procedures (Bown and Young, 1998). No physical concentration or separation was applied in order to retain the original assemblage composition. A polarizing light microscope at 1250× magnification was used to investigate the nannofossil assemblages through, at least, a longitudinal traverse of the smear slide (200 fields of view). Also, a fraction of each sample was disaggregated and washed through sieves, the residue being separated into three fractions (>200 µm, 100–200 µm and 50–100 µm). For each sample, the two great sized residues were investigated for planktonic foraminiferal content, although the richest assemblages were found in the 100–200 µm residues. A portion of the selected samples was separated for geochemical analyses, including carbon isotope, elemental and biomarker characterization.

#### 3.2. Oxygen and Carbon isotopes

Oxygen and carbon stable isotope analyses were performed on marls collected from the four sections, and also on carbonates from

the Puentenansa section. Carbonero and La Frontera samples were powdered using an agate mill. The  $\delta^{13}\text{C}$  values of carbonates were determined by treatment with 103% orthophosphoric acid using a VG Isocarb system thermostated at 90 °C. The produced  $\text{CO}_2$  was analyzed with an IRMS (Isotope Ratio Mass Spectrometer) VG Prism II. For analysis of organic matter  $\delta^{13}\text{C}$  values, samples were treated with 3% HCl for 24 h to remove carbonates and then analyzed with an elemental analyzer Carlo Erba 1108 coupled to a IRMS VG Isochrom in continuous flow mode. These analyses were performed at the Stable Isotope Laboratory (SIDI) of the Universidad Autónoma de Madrid and Servicio General de Isótopos Estables of the University of Salamanca (Spain). The results are expressed in the common  $\delta$ -notation in per mil (‰) relative to the VPDB-standard. The international carbonate standard NBS-19 (National Bureau of Standards;  $\delta^{13}\text{C} = 1.95\text{‰}$  and  $\delta^{18}\text{O} = -2.20\text{‰}$ ) was used to calibrate to V-PDB, with an average precision of 0.1‰ for  $\delta^{13}\text{C}_{\text{carb}}$  and 0.15‰ for  $\delta^{13}\text{C}_{\text{org}}$  analyses. A description of the method used for the Puentenansa and Cau sections has been published in Najarro et al. (2011b) and de Gea et al. (2003), respectively.

### 3.3. Elemental analyses and total organic carbon

A total of 33 samples from the studied sections were analyzed for total organic carbon (TOC) concentrations and elemental composition (CNHSO and XRF). The CNHSO and X-ray fluorescence analyses were performed in the Centro de Instrumentación Científica of the Universidad de Jaén, with a Thermo Finnigan FlashEA1112 CHNS-OElemental Analyzer, and an X-Ray spectrometer Bruker AXS model Pioneer S4 explorer, respectively. The TOC analyses were performed in the Centre of Atomic Spectrometry of the Universidad Complutense (Madrid) with a SHIMADZU TOC-V device that performs the 680 °C combustion catalytic oxidation method. The TOC concentration was determined by subtracting the inorganic carbon (IC) concentration from the total carbon (TC) in each sample.

### 3.4. Biomarkers

A collection of 15 samples, with an emphasis on the OAE1a intervals, were selected from the studied sections (Figs. 3 and 4), in order to characterize the thermal maturity and organic source inputs. The samples were externally washed with dichloromethane (DCM) to remove contamination from handling and then crushed using a Pulverisette 5 agate mill. The crushed samples were Soxhlet extracted for 48 h with 500 ml DCM/MeOH (80:20). Activated copper turnings were added to the solvent flask to remove elemental sulphur from the Carbonero samples. After extraction, the turnings were removed, and the solvent volume was reduced under reduced pressure. The extracts were separated into three fractions by column chromatography using activated alumina and sequential elution with hexane (saturated hydrocarbons), hexane:dichloromethane (9:1) (aromatic compounds), and methanol (polar compounds).

The saturated hydrocarbon fractions were analyzed with gas chromatography–mass spectrometry (GC–MS). GC–MS was carried out on a Thermo DSQ II gas chromatograph connected to a Thermo Trace Ultra mass spectrometer in the Centro de Instrumentación Científica of the Universidad de Jaén, and a ThermoQuest Trace GC–MS housed in the Organic Geochemistry Unit (School of Chemistry) of the University of Bristol. The GC was fitted with a fused silica capillary column (Supelco Equity–5; 30 m x 0.25 mm x 0.25  $\mu\text{m}$ ) and was operated with helium as carrier gas. The samples (in hexane) were injected at 70 °C and the oven was subsequently programmed to 130 °C at 20 °C/min and then at 4 °C/min to 300 °C where it was held for 25 min. Biomarkers were identified by comparison of mass spectra and retention time with those reported in the literature (i.e. Peters et al., 2005); for the identification of gammacerane an authentic standard purchased by Chiron was co-injected.

## 4. Results

### 4.1. Chemo- and biostratigraphy

The carbonate ( $\text{C}_{\text{carb}}$ ) and bulk organic carbon ( $\text{C}_{\text{org}}$ ) isotopic profiles are shown for each of the four sections (Figs. 3 and 4). Part of the data included in the curves comes from previous papers dealing with single sections (see de Gea et al., 2003 for the Cau section, and Najarro et al., 2011b for the Puentenansa section). The biostratigraphy of these successions is mostly based on the references cited and also on new data from this study. The reference curve used here for correlation is that published by Menegatti et al. (1998), because is the more widely used (i.e. Immenhauser et al., 2005; Föllmi et al., 2006; Dumitrescu and Brassell, 2006; Stein et al., 2011; Jenkyns et al., 2012), and it is focused on the same time interval studied here. Other more recent proposals of subdivision of the Barremian–Aptian C-isotope curve have also been considered (Bralower et al., 1999; Wissler et al., 2003; Herrle et al., 2004).

#### 4.1.1. La Frontera section

The  $\delta^{13}\text{C}_{\text{carb}}$  values of the La Frontera section range between 1.0‰ and 4.0‰ (Fig. 3). The lowest 9 samples in the section, corresponding to the latest Barremian marls and marly limestones of the Carretero Formation (*Micrantholithus hoschulzi* nannofossil biozone, Aguado et al., 1992a), show little isotopic variation, with values of  $\delta^{13}\text{C}_{\text{carb}}$  ranging from 1.8‰ to 2.1‰ (Fig. 3). A stratigraphic gap, including probably part of the uppermost Barremian and the lowermost Aptian, has been interpreted to exist at the top of this lower portion of the section, because the marker species *Hayesites irregularis* is not recorded in the uppermost part (uppermost Barremian) of the Carretero Formation and the overlying black facies of the Carbonero Formation are within the ‘nannoconid crisis’ interval (Fig. 3). This stratigraphic gap affects the complete lower member of the Carbonero Formation (lowermost Aptian), which is not represented in this section. A minor 0.5‰ positive excursion in carbonate, with a maximum value of 2.3‰ occurs within the middle member of the Carbonero Formation, in layers belonging to the *H. irregularis* nannofossil biozone and located within the interval corresponding to the ‘nannoconid crisis’ (Fig. 3). This excursion is followed by 10 m of largely carbonate-free black-shales, after which the positive excursion apparently continues to a maximum value of 4.0‰. *Eprolithus floralis* was recorded in the first sample above the carbonate-free rocks. Within the following 8 m interval, corresponding to the lowermost part of the *R. angustus* nannofossil biozone and *S. cabri* planktonic foraminiferal biozone, the  $\delta^{13}\text{C}_{\text{carb}}$  values remain relatively high (minimum 3.2‰; average 3.6‰). High abundances of *Assipetra*, (*A. infractetacea* and *A. terebrodentaria*), including large-sized specimens (*A. infractetacea* ssp. *larsonii* and *A. terebrodentaria* ssp. *youngii*; Tremolada and Erba, 2002), characterize this interval from the base of the middle member of the Carbonero Formation (Fig. 3). The next 5 m of the isotopic profile display an overall negative trend reaching a minimum of 1.0‰, coinciding with the *Globigerinelloides ferreolensis* and lower part of *G. algerianus* planktonic foraminiferal biozones (Lower–Upper Aptian transition). Finally,  $\delta^{13}\text{C}_{\text{carb}}$  values in the uppermost 12 m of the investigated interval are characterized by a slight increasing trend (maximum 2.8‰ but also a marked minimum value of 1.9‰). This last interval is Late Aptian in age, and is characterized biostratigraphically by the upper part of *G. algerianus* planktonic foraminifer biozone.

The  $\delta^{13}\text{C}_{\text{org}}$  values of the La Frontera section range between −21.0‰ and −27.4‰, and present a continuous record of C-isotope data. Generally, the  $\delta^{13}\text{C}_{\text{org}}$  values parallel the  $\delta^{13}\text{C}_{\text{carb}}$  curve. In detail, a large negative excursion, from −23.9‰ to −27.4‰, occurs in the interval lacking a  $\delta^{13}\text{C}_{\text{carb}}$  record. The subsequent positive excursion is characterized by an increase in  $\delta^{13}\text{C}_{\text{org}}$  values to −23.8‰ and this does parallel the  $\delta^{13}\text{C}_{\text{carb}}$  profile. However, also present in the upper part of the



$\delta^{13}\text{C}_{\text{org}}$  profile are several high frequency and high amplitude variations, with maximum shifts of 4.0‰ (Fig. 3).

#### 4.1.2. Carbonero section

The samples from the Carbonero section were taken in the type area of the Carbonero Formation (de Gea et al., 2008a). In this section, a stratigraphic gap has been interpreted to exist between the uppermost Barremian marls and marly limestones of the Carretero Formation and the overlying bluish-grey marls with intercalations of calcareous turbidites comprising the lower member of the Carbonero Formation (Lower Aptian). This gap, with a lower extent than that observed in the La Frontera section, has been deduced on the basis of the absence of the nannofossil *H. irregularis* in the Carretero Formation. The bluish-grey marls of the lower member of the Carbonero Formation, containing moderate to high proportions of nannoconids and *H. irregularis*, predate the 'nannoconid crisis' event (de Gea et al., 2008a). The base of the overlying black shales and radiolarian marls of the middle member of the Carbonero Formation (local expression of the OAE 1a), coincides with the onset of the 'nannoconid crisis'. For this reason, a new stratigraphic gap has been interpreted to exist in this section between the lower and middle members of the Carbonero Formation (de Gea et al., 2008a). Finally, the FO of the nannofossil *E. floralis*, has been recorded near the base of the alternation of greenish-grey marls and calcarenites of the upper member of the Carbonero Formation.

The Carbonero section has been analyzed at a lower resolution than the La Frontera section, but both the  $\delta^{13}\text{C}_{\text{carb}}$  and  $\delta^{13}\text{C}_{\text{org}}$  profiles are complete and largely parallel each other. Overall, they exhibit the same general trends as observed in the La Frontera Section (Fig. 3).  $\delta^{13}\text{C}_{\text{carb}}$  values vary between 3.1‰ and -7.9‰. The lower 20 m of the section (lower member of the Carbonero Formation), have constant values around 0.8‰ but then increase slightly up to 1.2‰. The following 28 m is characterized by a dramatic negative trend to values of approximately -7.9‰ in the radiolarian marls of the middle member of the Carbonero Formation. The next 15 m record a positive trend, up to 1.5‰, and the upper 80 m are characterized by generally elevated values (from 1 to 3‰).

The  $\delta^{13}\text{C}_{\text{org}}$  profile reproduces some of the major trends recorded by  $\delta^{13}\text{C}_{\text{carb}}$  values, albeit at a reduced amplitude. Values range from -28.9‰ to -24.4‰, although only a 2‰ decrease occurs from 30 to 40 m, such that the magnitude of the negative carbon isotope excursion is much smaller than that recorded by carbonates.

#### 4.1.3. Cau section

The studied interval of the Cau section belongs to the Almadich Formation. The lower member of the Almadich Formation in the Cau section encompass the Upper Barremian to lowermost Aptian interval and consists of a rhythmic alternation of marly limestones and light grey marls. The FO of *H. irregularis* has been recorded immediately above a ferruginous surface located within this lower member. As the levels below the ferruginous surface were assigned to the *M. hoschulzii* nannofossil biozone, a stratigraphic gap, probably affecting the uppermost Barremian and the lowermost Aptian, was interpreted to exist within this lower member (Aguado et al., 1999). The onset of the 'nannoconid crisis' event and an increase in the abundance of large-sized morphotypes of the nannofossil genus *Assipetra* (*A. infractacea* ssp. *larsonii* and *A. terebrodentaria* ssp. *youngii*; Tremolada and Erba, 2002) were observed within this lower member about 15 m above the ferruginous surface (Aguado et al., 1999). The middle member of the Almadich Formation is mainly composed of black shales and marls and its base coincides with the FO of the planktonic foraminifer *Schackoina cabri*. The FO of *E. floralis* has been recorded near the base of the marly limestones and grey marls characterizing the upper member of the Almadich Formation.

The  $\delta^{13}\text{C}_{\text{carb}}$  profile shows a clear pattern, with the same general trends observed in the La Frontera and Carbonero sections, with

two prominent positive excursions separated by a negative shift. The lowest part of the curve is poorly documented, as it is represented by only two analyses, with a lowest value of 1.4‰, and is capped by a discontinuity at the Barremian-Aptian boundary. The first positive excursion, with a maximum value of 2.7‰, is recorded within the *Deshayesites weissi* ammonite biozone (Fig. 4), predating the onset of the 'nannoconid crisis'. The subsequent negative excursion attains a minimum value of 0.4‰ in the uppermost part of the *H. irregularis* calcareous nannofossil biozone (equivalent to *G. blowi* planktonic foraminiferal biozone). The upper positive excursion, occurring in the *Deshayesites deshayesi* and *Dufrenoyia furcata* ammonite biozones, spans over 50 m and achieves a maximum value of 3.7‰ (Fig. 4). The  $\delta^{13}\text{C}_{\text{org}}$  record parallels that of the  $\delta^{13}\text{C}_{\text{carb}}$ , but with a lower sharpness (Fig. 4). de Gea et al. (2003) proposed a C-isotope stratigraphy, correlated with biostratigraphy published in Aguado et al. (1999), in which they recognized the eight segments first proposed by Menegatti et al. (1998) (Fig. 4). Although the C-isotope curve does not have a high-resolution, and some uncertainty in the segment boundaries can be acknowledged, the C3-C4 boundary, in the uppermost part of *B. blowi* biozone, is located at the base of the organic-rich interval, which records the main positive excursion.

#### 4.1.4. Puentenansa section

The studied interval in the Puentenansa section corresponds to the Patrocinio Formation, which consists, in this section, of a 31 m thick succession of open marine argillaceous and silty marls overlying a 13 m thick unit of rudist, gastropod and coral limestones and orbitolinid marls. The contact between the two units is an unconformity represented by a dissolution surface coated by a thin ferruginous crust, suggesting a brief episode of emersion at the top of the limestone followed by a flooding surface. Above this surface, deposition of the marls of the Patrocinio Formation occurred as consequence of platform drowning as the result of a combined action of a relative sea-level rise and poisoning by siliciclastic (Najarro et al., 2011a). According to the study of nannofossil assemblages in the studied section, the first 22 m of the Patrocinio Formation belong to the upper half of the *H. irregularis* Zone, and the absence of narrow canal nannoconids has allowed the identification of the 'nannoconid crisis' within this interval (Najarro et al., 2011b). The FO of *E. floralis* is registered 5 m above the sample PN-9 (Fig. 4). This biostratigraphic event allows the assignment of the upper part of the Patrocinio Formation in the Puentenansa section to the *R. angustus* Zone, suggesting the existence of a stratigraphic gap affecting to the upper part of the Lower Aptian (Najarro et al., 2011b).

In this section the  $\delta^{13}\text{C}_{\text{carb}}$  record can be subdivided into three successive intervals (Najarro et al., 2011a), (Fig. 4). The basal interval coincides with the Rábago and Umbrera Formations. It shows relatively homogeneous and positive  $\delta^{13}\text{C}$  values in the Rábago Formation (mean of +2.2‰), and a significant and progressive decrease of about 1‰ through the Umbrera Formation (but still with positive values). The second interval of the  $\delta^{13}\text{C}$  curve correlates with the Patrocinio Formation. This interval is characterized by a notable negative excursion from values of -0.4‰ at the base of the interval to -4.5‰ at the top. This negative excursion is not gradual but punctuated by three negative peaks (-2.9‰, -4.1‰ and -4.5‰ respectively; Fig. 4). Finally, the third interval in the  $\delta^{13}\text{C}_{\text{carb}}$  record shows the return to positive values, corresponding to the Reocin Formation. The carbon isotope composition of the bulk organic matter measured across the Patrocinio Formation in the Puentenansa section ( $\delta^{13}\text{C}_{\text{org}}$  in Fig. 4) ranges between -21.2‰ and -25.2‰, and shows three prominent negative spikes. From the bottom, the C-isotope curve starts with values of -22.2‰ and decreases sharply to -24.8‰, resulting in a first negative excursion of 2.6‰ in magnitude. This is followed by a return to more positive values (-21.6‰). The subsequent  $\delta^{13}\text{C}_{\text{org}}$  values show a gradual decrease towards lower values, reaching a minimum of -25.2‰ (second negative spike of 3.5‰ in

magnitude). Then, the values become progressively higher up to values of  $-22.6\%$ . Finally, at the top of the Patrocinio Formation, the profile displays a third negative spike of  $-24.6\%$  ( $\sim 2\%$  in magnitude) above the hiatal unconformity (Fig. 4).

The  $\delta^{13}\text{C}_{\text{carb}}$  and  $\delta^{13}\text{C}_{\text{org}}$  profiles lack the C4 to C7 isotopic segments of Menegatti et al. (1998), due to a discontinuity (Fig. 4) (Najarro et al., 2011b). Therefore, in the Puentenansa section only segment C3 of the OAE1a was recorded (Fig. 4). This section differs from the sections studied in the SIPM mainly due to the presence of shallow water carbonates at the base of the Aptian and in the Upper Aptian (Fig. 4).

#### 4.2. Elemental geochemistry

The TOC contents across all four sections vary between 0.5% and 5.6% (Table 1). Higher TOC contents occur in Carbonero and La Frontera sediments (average TOC = 3.8% and 1.9%, respectively), whereas Cau and Puentenansa sediments have lower TOC contents (average TOC = 1.0% and 0.6% respectively). Sulphur is absent or present in only trace concentrations in La Frontera and Cau, whereas it is present in low concentrations in the other sections (Puentenansa average S = 0.1% and Carbonero average S = 0.22%). Major element distributions (Table 1) clearly differentiate the four sections: Cau samples are marls and Puentenansa are mudstones, whereas Carbonero and La Frontera samples present a higher variability related to the facies differentiation between marls, mudstones and radiolarian marls.

#### 4.3. Distributions of hydrocarbon biomarkers

The extractable hydrocarbons are dominated primarily by *n*-alkanes, acyclic isoprenoids, steranes and hopanes (Fig. 5), although some sections also contain strong low-molecular-weight (Cau) or high-molecular-weight (Carbonero) unresolved complex mixtures (UCM). *n*-Alkanes are saturated straight-chain compounds (Fig. 5), mainly derived from vascular plants, both from aquatic (short-chain) and terrestrial (long-chain) environments (e.g. Meyers, 1997); however, attributing a specific source to any *n*-alkanes in the more thermally mature settings (see below) is problematic due to catagenetic alterations. Acyclic isoprenoids are composed of polymerized isoprene units, and the dominant compounds identified here, pristane and phytane, are typically considered to be derived from chlorophyll (e.g. Powell and McKirdy, 1973; but see ten Haven et al., 1987). Hopanes are  $\text{C}_{27}$ – $\text{C}_{35}$  pentacyclic triterpenoids and originate from bacterial bacteriohopanepolyols (Ourisson et al., 1982). Steranes are tetracyclic triterpanes arising from diagenetic alteration of sterols and consequently almost exclusively derive from eukaryotic organisms (e.g. Moldovan et al., 1985). In most samples, the *n*-alkanes, acyclic isoprenoids and hopanes dominate, with variable ratios of *n*-alkanes/hopanes, and steroids represent a lower contribution (Fig. 5).

##### 4.3.1. *n*-Alkanes

The *n*-alkanes are present and typically the dominant compounds in most chromatograms. Short and long-chain *n*-alkanes are present, ranging from  $n\text{-C}_{13}$  up to  $n\text{-C}_{35}$  (Fig. 5), in all samples, but the distribution varies significantly among the samples studied, both between different sections and stratigraphically. In the Puentenansa and La Frontera sections there is a clear and strong bimodal distribution, represented by low-molecular-weight (LMW;  $<\text{C}_{22}$ ) components with no carbon preference and high-molecular-weight-components (HMW;  $>\text{C}_{24}$ ) that have an odd-over-even carbon number predominance (OEP) at La Frontera but not at Puentenansa (Fig. 6). In contrast, all of the Carbonero rocks samples are dominated by short-chain compounds and the HMW components have no carbon number preference. Somewhat intermediate distributions occur in the Cau samples.

**Table 1**  
TOC, and elemental composition of selected samples.

Sample	Thickness (m)	TOC %	N% %	C% %	S% %	Al <sub>2</sub> O <sub>3</sub> %	SiO <sub>2</sub> %	CaO%
<b>La Frontera</b>								
XF-103.1	43.25	1.1	0.0467	3.9193	0			
XF-96.2	26.90	1.4	0.0478	3.8031	0			
XF-1-33	25.7	0.7	0.0612	0.5714	0			
XF-1-27	24.5	0.8	0.063	0.6515	0			
XF-1-23	23.4	1.7	0.0945	1.3781	0.0397			
XF-1-20						13.89	59.19	0.78
XF-1-13						13.26	58.35	1.04
XF-1-1						9.47	29.8	23.98
XF-90.16	22.20	4.5	0.0296	6.7333	0	6.05	26.35	29.83
XF-90.5	20.00	0.7	0.0318	6.0533	0	7.49	27.24	27.61
XF-79	13.50	4.1	0.0418	5.517	0			
XF-72	9.30	4.8	0.0683	7.2018	0			
<b>Puentenansa</b>								
PN-14	29	0.5	0.0513	0.1489	0.0014	20.89	46.66	0.33
PN-13	27	0.8	0.0439	0.3525	0.0112	20.02	46.83	0.84
PN-10	21	0.6	0.0473	0.6973	0.0092	19.81	48.54	1.81
PN-7	15	0.6	0.0431	0.5447	0.2433			
PN-5	11	0.9	0.0418	1.1211	0.1706			
PN-4	9	0.7	0.0443	0.7718	0.1178			
PN-3	7	0.4	0.0275	3.7348	0.0123			
PN-2	5	0.6	0.0425	1.216	0.2702			
PN-1	3	0.5	0.0436	1.4614	0.292			
<b>Cau</b>								
Cau-32a	88.3	0.9				8.14	16.55	37.12
Cau28a	60.6	0.7	0.0225	9.8446	0	5.57	11.3	42.69
Cau23a	52.5	1.9	0.0338	9.3192	0.006	7.09	14.24	39.12
Cau20a	46	1	0.0305	9.4636	0.0071			
Cau19a	42	0.9	0.0279	9.2777	0.01	5.48	12.5	41.06
Cau16a	37.1	1	0.0337	8.6089	0.0369	6.98	16.84	36.76
Cau13a	33.6	0.8	0.0213	8.2714	0.0138	7.57	17.99	35.94
Cau12a	29.5	1.1	0.1051	0.9445	0.0846			
Cau10a	25.1	0.9	0.0192	9.2012	0			
<b>Carbonero</b>								
CAB-7	63.6	4.2	0.15	5.8234	0.1344			
CAB-6	53	5.6	0.18	5.3984	0.1254			
CAB-5	51	4.8	0.14	5.3118	0.7261	9.8	52.16	8.66
CAB-4	51	1.1	0.08	0.9067	0.2114	13.46	58.56	2.5
CAB-2C	40	3.7	0.1	6.6046	0.2951	10.3	35.05	19.71
CAB-2A	30.3	4.2	0.09	3.5869	0.0303	12.4	42.06	12.13
CAB-1J	25	3	0.0579	4.8499	0.0514	10.2	33.96	20.63

To facilitate interpretation of these *n*-alkane distributions, we have calculated two different ratios (Fig. 6): the HMW/LMW ratio ( $[\text{n-C}_{25} + \text{n-C}_{26} + \text{n-C}_{27} + \text{n-C}_{28} + \text{n-C}_{29}] / [\text{n-C}_{17} + \text{n-C}_{18} + \text{n-C}_{19} + \text{n-C}_{20} + \text{n-C}_{21}]$ ) in order to constrain the relationship between long and short-chain compounds, and also the odd-over-even predominance ratio ( $\text{OEP1} = \text{C}_{21} + (6 \times \text{C}_{23}) + \text{C}_{25} / (4 \times \text{C}_{22}) + (4 \times \text{C}_{24})$ , and  $\text{OEP2} = \text{C}_{25} + (6 \times \text{C}_{27}) + \text{C}_{29} / (4 \times \text{C}_{26}) + (4 \times \text{C}_{28})$ ), defined by Scalan and Smith (1970). We note that the HMW/LMW ratio has also been presented as the Terrestrial to Aquatic Ratio (TAR; Bourbonniere and Meyers, 1996), but the TAR terminology is not used in this context because of the prominent control of thermal maturity on calculated ratios (see below). There is a clear OEP (Table 2 and Fig. 6) in all of the La Frontera and Cau samples ( $\text{OEP1} = 1.07\text{--}1.73$ ;  $\text{OEP2} = 1.75\text{--}3.49$ ), but it is low or absent in the Puentenansa and Carbonero samples (PN:  $\text{OEP1} = 1.37\text{--}1.53$ ;  $\text{OEP2} = 0.60\text{--}1.59$ ; CAB:  $\text{OEP1} = 0.71\text{--}1.35$ ;  $\text{OEP2} = 0.84\text{--}1.41$ ; Table 2). Similarly, all Carbonero samples are characterized by a dominance of short-chain *n*-alkanes (average HMW/LMW ratios of  $0.62 \pm 0.36$ ). HMW/LMW ratios at Puentenansa are similar to those at Carbonero ( $0.69 \pm 0.47$ ), but the distribution is different, as it is bimodal (see Fig. 5). In the other sections, the HMW *n*-alkanes are proportionally more abundant (HMW/LMW ratios at La Frontera =  $0.83 \pm 0.61$  and Cau =  $1.33 \pm 0.17$ ). With respect to vertical trends, we acknowledge that we have limited data from each section; nonetheless, the CAB and CAU sections, with



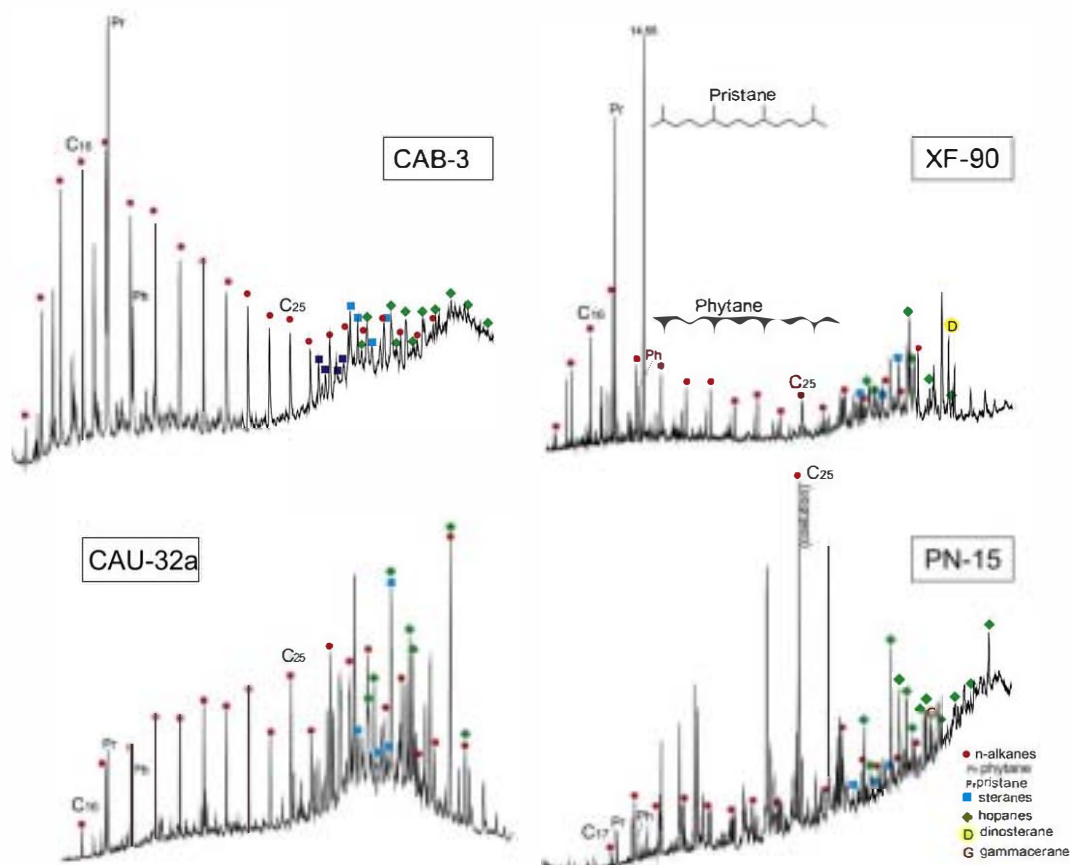


Fig. 5. Selected total ion chromatograms of samples from the studied sections. Numbers refer to *n*-alkanes.

greater sample resolution, show a clear variability through the section in the HMW/LMW ratio (Fig. 6).

#### 4.3.2. Isoprenoids

Pristane and phytane are abundant in all samples and exhibit high variability in their relative abundances. The Pr/Ph and isoprenoid/*n*-alkane ratios vary strongly, both between different sections and stratigraphically (Table 2 and Fig. 6). Pr/Ph ratios are highest in the Carbonero (Pr/Ph = 2–4.65, average = 2.98) and La Frontera samples (Pr/Ph = 2.08–4.6, average = 3.4), have intermediate values in the Cau samples (0.67–3.15, average = 1.64) and have the lowest values in the Puentenansa samples (0.6–0.68). At the Cau and Carbonero sections where a more complete suite of samples has been analyzed, Pr/Ph ratios generally decrease upsection, with highest values coinciding with pre-negative  $\delta^{13}\text{C}$  excursion levels. The isoprenoid/*n*-alkane ratios (Fig. 6) are also highly variable, with generally higher Pr/*n*-C<sub>17</sub> ratios and lower Ph/*n*-C<sub>18</sub> ratios in the Carbonero Section than at Cau (Table 2 and Fig. 6).

#### 4.3.3. Hopanes

Hopanes are present in all of the samples studied, ranging in carbon number from C<sub>27</sub> to C<sub>35</sub>. The C<sub>30</sub> component is dominant in the Carbonero and La Frontera samples, C<sub>31</sub> is dominant or equivalent to C<sub>30</sub> in Cau samples, and the C<sub>29</sub> hopane is dominant in the Puentenansa section (Fig. 7). The C<sub>31</sub>–C<sub>35</sub> extended hopanes (homohopanes) are present in CAB and PN samples, whereas in CAU and XF samples only C<sub>31</sub>–C<sub>32</sub> compounds are present. The commonly observed decrease in hopane concentration from C<sub>31</sub> to C<sub>35</sub> occurs in all samples, except for PN-15, where the C<sub>35</sub> component is somewhat more abundant than the C<sub>34</sub>. Crucially, the distribution of hopane stereoisomers differs dramatically among the four sections (Fig. 7 and Table 3). The 22S/(22S + 22R) homohopane ratio is 0.5

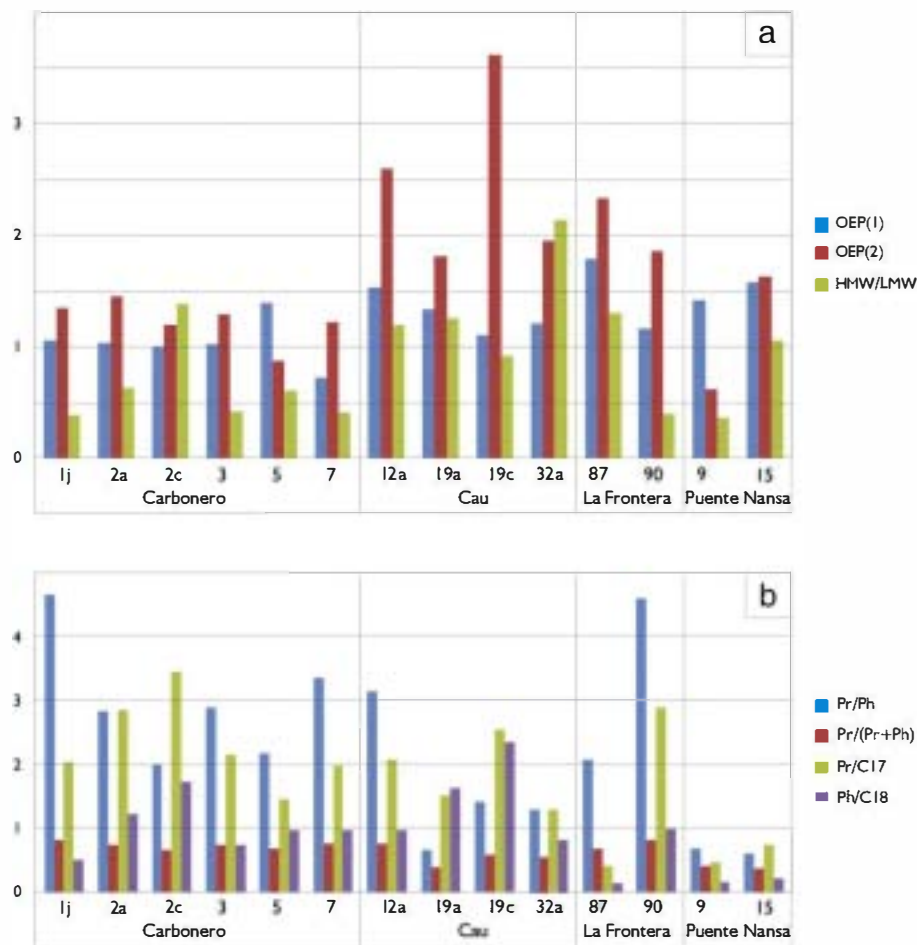
in the Carbonero and Puentenansa samples, whereas in La Frontera and Cau this ratio is 0.13–0.37. Similarly, the 17 $\alpha$ ,21 $\beta$ (H) isomers are dominant in the Carbonero and PN sections, with subordinate abundances of the 17 $\beta$ ,21 $\alpha$ (H) diastereomers (moretanes), whereas the Cau samples are characterized by a dominance of hopane diastereomers with the 17 $\beta$ ,21 $\beta$ (H) configuration, although 17 $\alpha$ ,21 $\beta$ (H) and 17 $\beta$ ,21 $\alpha$ (H) diastereomers are also present. The Ts and Tm C<sub>27</sub> hopane diastereomers are present in all samples, with a Tm/Ts ratio  $\geq 1$  in the Carbonero, La Frontera and Cau samples and a Tm/Ts ratio <0.4 in Puentenansa samples.

Gammacerane is present only in the Puentenansa samples (confirmed with standard co-injection), where this compound has a relatively high abundance in comparison with hopanes (G/C<sub>30</sub> = 0.58).

#### 4.3.4. Steranes

A range of C<sub>27</sub> to C<sub>30</sub> steranes occur in all of the sections studied (Fig. 8); in a broad sense, the relative distributions are similar, with C<sub>29</sub> always being dominant (48%–70%), C<sub>27</sub> intermediate (19%–28%), and C<sub>28</sub> minor (9%–31%), but considerable variability within that framework is present. Further variability is reflected by the abundance of a C<sub>30</sub> 4-methylsterane, tentatively interpreted to be dinosterane on the basis of its mass spectrum and retention time (Summons et al., 1987), which is present in some XF and CAU samples but is absent or occurs in only trace abundances in the PN and Carbonero samples. In fact, in one sample (Cau-24, see Fig. 8), it is one of the dominant compounds in the apolar fraction and certainly the most abundant sterane.

The sections also differ with respect to the relative abundances of sterane diastereomers. The CAU and XF samples are represented almost exclusively by the 5 $\alpha$ (H),14 $\alpha$ (H),17 $\alpha$ (H),20R epimer. On the other hand, in CAB and PN samples, 5 $\alpha$ (H),14 $\beta$ (H),17 $\beta$ (H) diastereomers and 22S steranes are also relatively abundant, with similar



**Fig. 6.** Biomarker ratios derived from relative *n*-alkane (a) and isoprenoid (b) abundances. Ratio definitions and interpretations are provided in the text. Location of the samples shown in Figs. 3 and 4.

relative distributions in both sections (Fig. 8). Furthermore, the ratios of the 20S and 20R epimers differ between the latter two sections, with the 20S/20R ratio being around 1 for the 5 $\alpha$ (H),14 $\alpha$ (H),17 $\alpha$ (H) diastereomer in CAB samples, whereas in PN samples the

20R stereochemistry is dominant. Similarly, 13 $\beta$ ,17 $\alpha$  steranes (diasteranes) are relatively abundant compared to regular steranes in all of the Carbonero samples but absent in the other sections.

## 5. Discussion

### 5.1. Correlation of C-isotope curves and integration with biostratigraphic data and distribution of organic-rich facies

To correlate the four sections, we have used the biostratigraphic data and also attempted to identify the eight segments proposed by Menegatti et al. (1998) for the Aptian C-isotope profile from the “Selli level” in the Alps. Those authors proposed a subdivision of the Aptian C-isotopic record in 8 segments based on changes observed in two different sections in the Alpine domain and attempted to correlate those carbon isotopic temporal variations with nannofossils and planktonic foraminifers biozonations (Fig. 9). The Cau and La Frontera sections both contain the eight segments described by Menegatti et al. (1998), whereas in the Carbonero section there are no isotope data for the C1 segment, although biostratigraphic data suggest it is present. In Puenteansa, the succession starts within the C2 segment, and there is a discontinuity affecting the C4 to C7 segments (see Figs. 3, 4 and 9).

The sections exhibit the expected correlations between the biostratigraphic data and the C-isotope curves (i.e. segments proposed by Menegatti et al., 1998), and consistent relationships are observed in all four Iberian sections studied with the exception of the FO of *E. floralis*. This datum was recorded within the C6 segment in La

**Table 2**  
n-Alkanes and isoprenoids ratios.

Sample	Pr/Ph	Pr/(Pr+Ph)	Pr/C17	Ph/C18	OEP (1)	OEP (2)	HMW/LMW
<b>Carbonero</b>							
CAB-1J	4.65	0.82	2.05	0.51	1.02	1.30	0.38
CAB-2A	2.84	0.74	2.86	1.23	1.00	1.41	0.62
CAB-2C	2.00	0.67	3.46	1.73	0.98	1.17	1.34
CAB-3	2.89	0.74	2.15	0.75	0.99	1.25	0.42
CAB-5	2.18	0.69	1.45	0.97	1.35	0.84	0.60
CAB-7	3.36	0.77	1.98	0.98	0.71	1.18	0.34
<b>Cau</b>							
CAU-12a	3.15	0.76	2.07	0.97	1.49	2.51	1.16
CAU-19a	0.67	0.40	1.52	1.63	1.29	1.75	1.22
CAU-19C	1.42	0.59	2.54	2.34	1.07	3.49	0.90
CAU- 32 a	1.32	0.57	1.30	0.84	1.17	1.88	2.06
<b>La Frontera</b>							
XF-87	2.08	0.68	0.41	0.14	1.73	2.25	1.26
XF-90	4.60	0.82	2.90	0.99	1.12	1.80	0.39
<b>Puenteansa</b>							
PN-9	0.68	0.41	0.46	0.16	1.37	0.60	0.36
PN-15	0.60	0.38	0.75	0.22	1.53	3.00	1.02



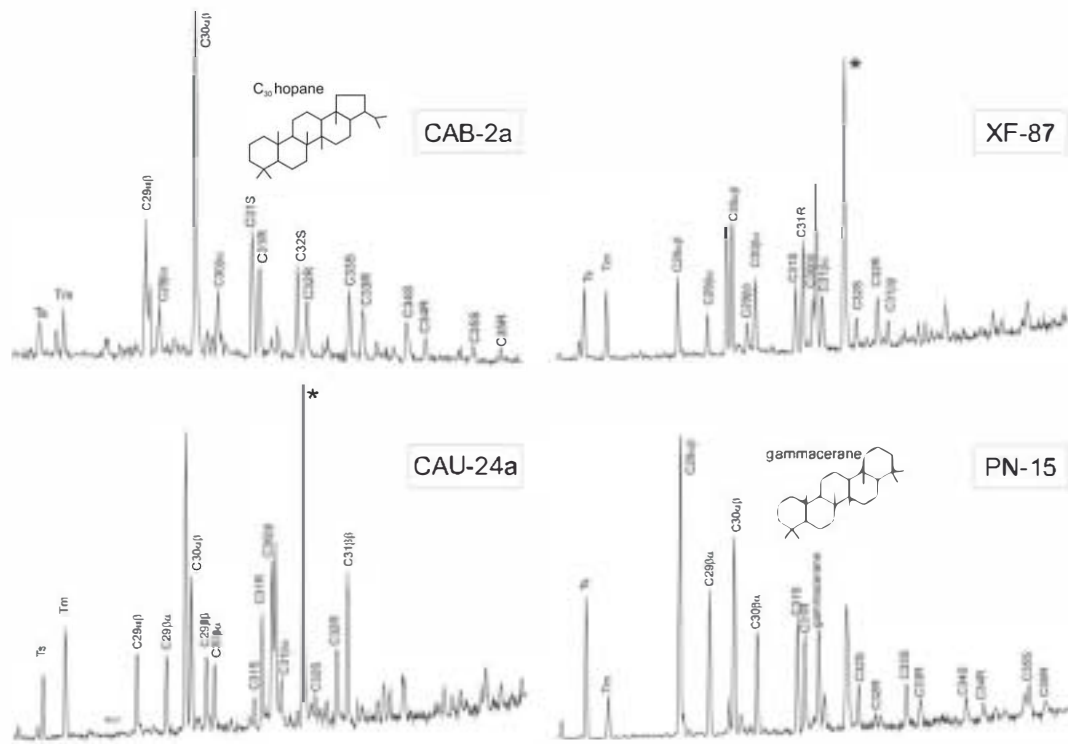


Fig. 7. Selected partial mass ( $m/z = 191$ ) chromatograms of samples from the studied sections, showing hopanes and gammacerane (\* unknown compound, probable ketone or diahopane).

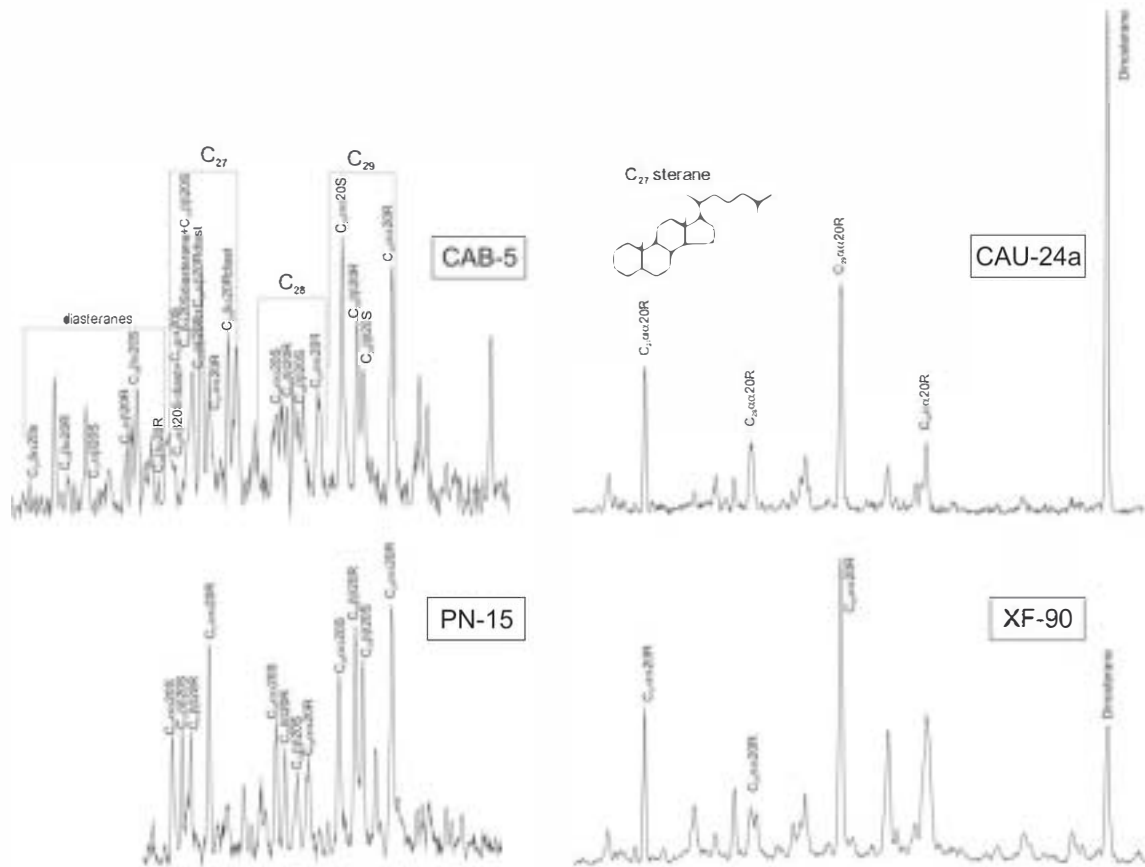
Frontera section, and within the C7 segment in Carbonero and Cau sections. We suspect that the diachronism in the record of the FO of *E. floralis* in Carbonero section is related to the poor preservation of the nannofossil assemblages in the middle member and lowest part of the upper member of the Carbonero Formation. In most of the samples from this middle member, nannofossils were not found or were very poorly preserved.

In the Cau section, however, the cause of this diachronism cannot be attributed to preservation, as nannofossil assemblages are diverse and moderately well preserved in this interval, and we suspect that it is probably related to palaeoecological factors. The palaeoceanographic setting of the Cau section was more proximal and shallower (outer platform) than that of the La Frontera section (pelagic). In addition, the

nannofossil assemblages from this interval in the Cau section contain higher proportions of nannoconids and pentaliths (*Braarudosphaera*, *Micrantholithus*) than those of the La Frontera section. Nannoconids are well-documented Cretaceous taxa predominantly associated to low-latitude carbonate-shelfal and epicontinental basins likely with stratified waters (Thierstein, 1976; Roth and Bowdler, 1981; Roth and Krumbach, 1986; Busson and Noël, 1991; Mutterlose, 1992; Street and Bown, 2000; Bown, 2005). They are also related to oligotrophic surface waters in which salinity may also have played a significant control (Lees et al., 2005). Erba (1994) suggested that nannoconids were analogous to the extant genus *Florisphaera*, which proliferates in the deep photic zone of modern oceans at a deep nutricline. This author suggested that the OAE1a-related nannoconid crisis was the result of nutrification of surface waters, triggering blooms of surface waters coccolithophorids and depletion of deeper-dwelling nannoconids. Pentaliths (*Micrantholithus* and *Braarudosphaera*) appear to be similarly neritic to nannoconids in distribution. Its ecology is also probably related to neritic factors such as reduced salinity and enhanced nutrient content (Roth and Bowdler, 1981; Parker et al., 1985; Siesser et al., 1992; Street and Bown, 2000; Bown, 2005). *Micrantholithus* was replaced by *Braarudosphaera* in the Aptian, which appears to have retained a similar ecology from Aptian to recent times, being most common in neritic environments (Gran and Braarud, 1935). Other anomalous occurrences of *Braarudosphaera* blooms, linked to post-extinction assemblages (Cretaceous–Tertiary boundary) or to subtropical open-ocean sites (Parker et al., 1985; Scarparo-Cunha and Shimabukuro, 1997) suggest an opportunistic behavior for this taxon (Siesser et al., 1992). Given the paleoceanographic setting of the Cau section (distal part of a platform) and the above-mentioned higher abundances of ecologically significant nannoconids and pentaliths through the C6 and C7 carbon-isotope intervals, we suggest that a shallower, more restricted environment, with lower salinity, could be related with the delay in the first record of *E. floralis* in this section. Similar diachronisms in the FOs and LOs of several micro- and nannofossil indicators with respect to the carbon-isotope stratigraphy have been also detected for sections in the Mazagan Plateau and the Voconian Trough (Herrle et al., 2004), and

Table 3  
Maturity-related biomarker parameters (ratios of hopanes and steranes).

Hopanes			Steranes		
Sample	22S/ (22S + 22R) (C31)	C30 $\beta\alpha$ / ( $\beta\alpha$ + $\alpha\beta$ )	Sample	$\beta\beta/\beta\beta + \alpha\alpha$	20S/ (20S + 20R)
Carbonero			Carbonero		
CAB-7	0.53213368	0.14231738	CAB-7	0.302463054	0.516723549
CAB6a	0.54291845	0.11111111	CAB-5	0.34566787	0.4665195
CAB-5	0.54736842	0.13127413	CAB-3	0.44603033	0.432876712
CAB-3a	0.54174397	0.13378685	CAB-2C	0.399749373	0.505675955
CAB-2a	0.56915739	0.172322023	CAB-2A	0.371191136	0.512352309
CAB-1J	0.54363636	0.18592297	CAB-1J	0.331963001	0.492583919
La Frontera			Puentenansa		
XF-90	0.22988506	0.33174603	PN-15	0.466313763	0.434472208
XF-87	0.37260274	0.38235294			
Cau					
CAU-32A	0.131528046	0.581020591			
CAU-24a	0.21507761	0.32028986			
CAU-19a	0.225	0.32484076			
Puentenansa					
PN-15	0.52012384	0.32767402			
PN-9	0.50875146	0.42937063			

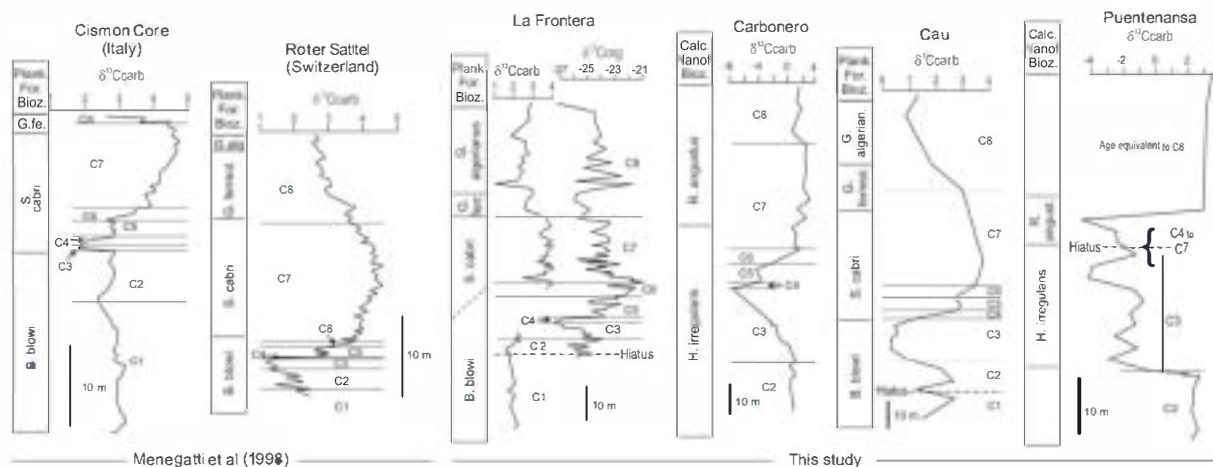


**Fig. 8.** Selected partial  $m/z = 217$  mass chromatograms of samples from the studied sections, showing steranes. Note that diasteranes are only present in CAB samples. Dinosterane, although present in these  $m/z = 217$  chromatograms, has a mass spectrum with a dominant 231 fragment and a minor 217 fragment.

interpreted in terms of regional differences in palaeoenvironmental conditions.

Taking this into account, the data presented in this study reveal several relationships, shown in [Figs. 3 and 4](#): (1) The C1–C2 boundary is affected by a discontinuity, at the Barremian–Aptian boundary, in all four sections; (2) the C3–C4 boundary corresponds to the base of *S. cabri* foraminifer biozone; (3) the base of the *R. angustus* nannofossil biozone is diachronic, and its first occurrence lies within the C6 segment; and (4) the base of the *G. ferreolensis* foraminifer biozone occurs within the C7 segment.

Correlation of biostratigraphy, C-isotope stratigraphy and facies among the four sections reveals a variety of spatial and temporal relationships ([Fig. 10](#)). Although C-isotope stratigraphy is correlatable around the world, the timing of deposition of organic-rich sediments is not coeval (i.e. [Jenkyns, 2010](#)), and that is apparent in the Iberian region of the Tethys. Organic-rich facies are first deposited in pelagic environments (La Frontera and Carbonero sections), at the base of the Aptian and corresponding to the C2 segment of the carbon isotope profile, whereas organic-rich black shales or marls are deposited during the C3 or C4 segments in platform settings (Puentenansa and Cau



**Fig. 9.** Correlation of the studied sections with the reference curves of [Menegatti et al., 1998](#).



## 5.2.1. Thermal maturity

The use of biomarker distributions to assess the thermal maturity of organic matter in sediments and rocks is well established (e.g. Peters et al., 2005). The most powerful tools at low to moderate maturity (with respect to oil generation) are those based on the relative distribution of sterane and hopane isomers, and that is the focus here. However, thermal maturity also affects *n*-alkane distributions and, therefore, informs our interpretation of between-section differences in those parameters.

The Carbonero section has 22S/(22S + 22R) homohopane ratios of ca 0.55, C30  $\beta\alpha/(\beta\alpha + \alpha\beta)$  hopane ratios of ca 0.15 (Fig. 11a),  $\beta\beta/(\beta\beta + \alpha\alpha)$  sterane ratios of 0.37, and 20S/(20S + 20R) sterane ratios of 0.49 (Table 3, Fig. 11b), and abundant diasteranes. All of these data indicate that OM from the Carbonero section is thermally mature with respect to oil generation (Mackenzie et al., 1980; Seifert and Moldowan, 1980; Waples and Machihara, 1991). Although the homohopane ratios are at a maximum and cannot put an upper limit on the section's thermal maturity, the persistence of these biomarkers—and especially the moretanes—suggests that the section has not exited the oil generation window. Sterane distributions also indicate that the section is near the peak stage of oil generation, equivalent to a vitrinite reflectance between 0.65 and 0.85 (according to Peters et al., 2005). The elevated thermal maturity accounts for the low HMW/LMW *n*-alkane ratios and the lack of an odd-over-even predominance.

The Cau and La Frontera sections, in contrast, are characterized by distributions consistent with low thermal maturity, including a dominance of 22R homohopanes (i.e. 22S/(22S + 22R) homohopane ratios of 0.13–0.37), the persistence of hopanes with the biological and thermally unstable 17 $\beta$ ,21 $\beta$ (H) configuration, steranes represented only by the  $\alpha\alpha$ 20R isomers, and an absence of diasteranes (Table 3, Figs. 7 and 8). These data indicate that Cau and La Frontera have very low maturity and explain why higher plant wax *n*-alkane distributions, including the high OEP and HMW/LMW ratios, have persisted. Puentenansa samples have hopane and sterane ratios indicating that the OM is thermally mature (within the peak stage of oil generation) but not to the same degree as the Carbonero samples; for example, diasteranes are not present in Puentenansa. Thus, differences in thermal maturity are a main driver of organic geochemical differences among the studied sections. Specifically, Carbonero and Puentenansa OM is thermally mature, whereas Cau and La Frontera OM is thermally immature, probably with an equivalent vitrinite reflectance lower than 0.4 (according to Peters et al., 2005). Particularly interesting are the important differences observed between Carbonero and La Frontera OM, taking into account that they were deposited in nearby areas. The higher degree of maturation showed in Carbonero samples could be related to the Cretaceous tectonic and volcanic activity occurred in the Carbonero area, related to the extensional tectonics (Molina et al., 1998; de Gea et al., 2008a, 2008b) which could have resulted in a heating of the sediments, whereas the nearby La Frontera area, not affected by these processes, records immature OM; however, a different burial history cannot be dismissed. The low thermal maturity in Cau and La Frontera sections suggests that functionalized biological compounds, including alcohols and carboxylic acids, will be preserved in these sections (which we have confirmed but will be discussed in subsequent papers). This will affect interpretation of OM sources and sedimentary environments and is an important caveat to subsequent sections.

## 5.2.2. OM sources and environmental conditions

Lipid biomarkers can be used to infer sources of organic matter (e.g. terrestrial vs marine) and depositional conditions (e.g. oxic vs. anoxic). Overall, all of the OM present in the studied samples is interpreted to derive from significant terrestrial inputs as well as

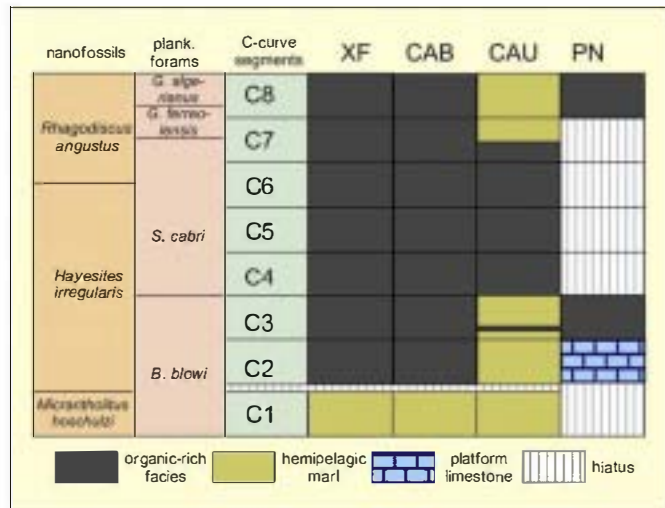


Fig. 10. Correlation between C-isotope stratigraphy (segments from Menegatti et al., 1998), biostratigraphy and facies.

sections, respectively). Also, the interval of deposition of black shales is longer in pelagic sections (up to the C8 segment) than in the platform sections (C7) in the SIPM. When comparing the palaeogeographically distinct platform settings (SIPM, Cau, and NCB, Puentenansa), the deposition of organic-rich facies starts earlier and persists longer in the North Cantabrian basin (C2 to C8) than in the South Iberian (C3 to “early” C7).

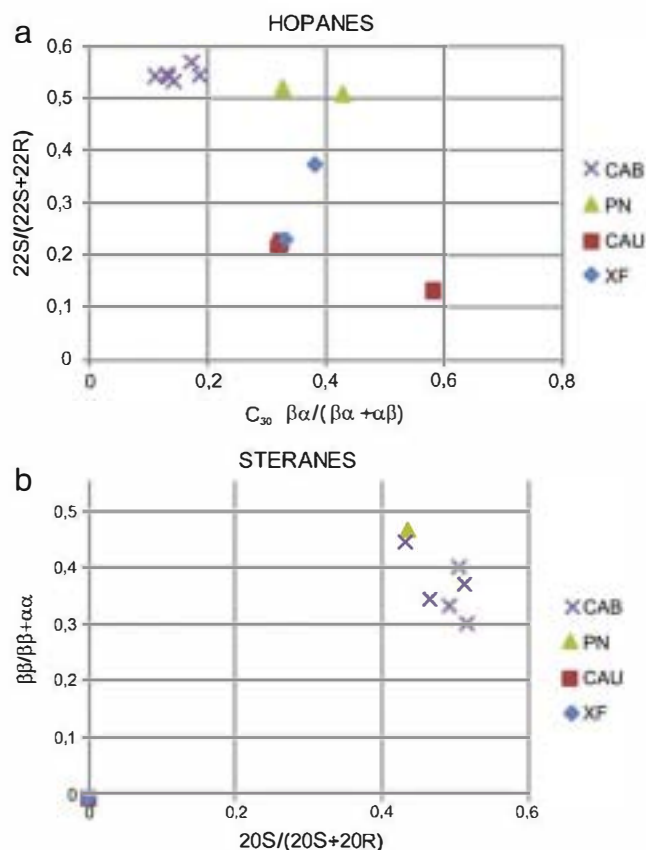


Fig. 11. Maturity-related biomarker parameters. a: 22S/(22S + 22R) homohopane ratios versus  $\beta\alpha/(\beta\alpha + \alpha\beta)$  hopane ratios. b:  $\beta\beta/(\beta\beta + \alpha\alpha)$  sterane ratios versus 20S/(20S + 20R) sterane ratios.

marine and bacterial sources. The differences in thermal maturity complicate comparison of biomarker parameters among the sections. Moreover, the high level of maturity in several samples prevents an accurate interpretation in terms of source. Nevertheless, biomarker data from this study do reveal some systematic relationships, allowing tentative comparisons.

**5.2.2.1. OM sources.** The predominance of high molecular weight *n*-alkanes ( $C_{25}$ – $C_{35}$ ) with an odd-over-even carbon number in the La Frontera and Cau sections indicate a terrestrial contribution from wax lipids of higher plants (Eglinton and Hamilton, 1967). The low molecular weight  $C_{14}$ – $C_{24}$  *n*-alkanes, as well as pristane and phytane, from the same sections likely derive from marine contributions. Puentenansa samples have a slight OEP, but still contain a strong HMW *n*-alkane ( $C_{25}$ – $C_{35}$ ) signal, which would also indicate a terrestrial contribution. The predominance of LMW *n*-alkanes in CAB samples reflects their thermal maturity rather than OM sources.

The steranes show a rather constant distribution with a dominance of  $C_{29}$  compounds in all samples. This feature could also indicate a strong terrestrial contribution (Huang and Meinschein, 1979), but caution is essential as many algae synthesize  $C_{29}$  sterols (Peters et al., 2005 and references therein). Further evidence that the high  $\%C_{29}$  steranes should be interpreted cautiously comes from the fact that there are no systematic differences between the platform and pelagic sections, even when comparing sections with similar thermal maturity. Marine inputs are clearly significant at all sites, despite the relatively low  $\%C_{27}$  and  $\%C_{28}$  values, indicated in particular by the occurrence of dinosterane derived from dinoflagellates (Withers, 1983). Dinosterane occurs in several samples in concentrations comparable to those of steranes (XF samples), is absent in others (CAB and PN samples, CAU-12, CAU-19a and CAU-19c), and is one of the most abundant hydrocarbons in CAU-24. Although the sample resolution is limited, there is strong variation in dinosterane concentrations within a given section, indicating that dinoflagellate production varied, perhaps reflecting changes in nutrient status. Moreover, the particularly high concentration in the CAU-24 sample suggests that these conditions arose when either upwelling or runoff delivered nutrients to the shallow marine environment. However, insufficient data prevent us from further interpretation of either temporal trends or differences between sections.

Bacterial sources of OM are demonstrated by the presence of hopanes in all samples. Also, although it is not a bacterial compound, gammacerane typically occurs when bacteria are present in the water column (Sinningh -Damst  et al., 1995; Peters et al., 2005). Intriguingly, the abundance of hopanes relative to *n*-alkanes is particularly high in samples CAB-5, XF-90, and CAU-19a, all corresponding to the interval located above the negative carbon isotope excursion. Previous work has shown that other times of global anoxia are associated with major changes in the bacterial population, and in particular the proportional contributions of cyanobacteria (e.g. Kuypers et al., 2004 -OAE2-, Xie et al., 2007 -Permian-Triassic Boundary), and future work will focus on similar changes in the Iberian region during OAE1a.

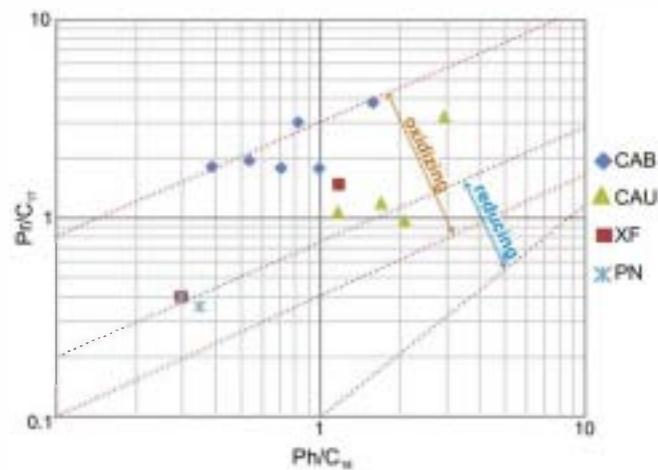
**5.2.2.2. Environmental conditions.** One of the main questions related to anoxic events is the geographical extent of reducing conditions in the water column. Gammacerane is present in the Puentenansa samples, and its principal source appears to be bacteriivorous ciliates, which occur at the interface between oxic and anoxic zones in stratified water columns (Sinningh -Damst  et al., 1995; Peters et al., 2005), such that its presence is often associated with anoxia induced by water column stratification. Intriguingly, we did not detect gammacerane in the Carbonero section, even though it is characterized by the highest TOC contents, and has sedimentologic evidence of anoxia (i.e. barite concretions, Molina and Hern ndez-Molina, 1993; de Gea et al., 2008a).

A range of other biomarker distribution based redox proxies have been developed (e.g. Peters et al., 2005), but these are based on

empirical observations of petroleum and corresponding source rocks rather than thermally immature sediments; thus, in our samples they can only be applied to the CAB and PN samples. The Pr/Ph ratio has been used to infer redox conditions, but is also governed by thermal maturity, source inputs and lithology (Didyck et al., 1978; Ten Haven et al., 1987; Hughes et al., 1995). Carbonero samples have a relatively high Pr/Ph ratio, between 2.3 and 5.5, whereas PN samples have very low values (0.4 and 0.5), with the latter being typical for anoxic conditions ( $<0.8$ ; Hughes et al., 1995). Other aspects of the PN biomarker profile are consistent with this, including the elevated concentrations of the  $C_{35}$ -homohopane (Peters and Moldowan, 1991) and the high gammacerane/hopane ratios. Collectively, all of the biomarker data suggest that CAB samples, despite being more organic-rich, reflect more oxygenated conditions than PN samples (Fig. 12). By extension, this suggests that more reducing environments prevailed in the North Cantabrian Basin than the SIPM.

### 5.3. Sedimentary model: integration of stratigraphy, geochemistry, palaeogeography and palaeoceanography

The differences observed in the timing of deposition of the organic-rich facies between sections can be explained in relation to the different palaeogeographic settings and the palaeoceanography (Fig. 13). During Menegatti's C2 interval, the organic-rich deposition only occurs in the deep pelagic sections of the SIPM (Carbonero and La Frontera), located in fault-bounded high-subsident areas, probably with reduced water circulation, whereas in the shallow marine settings normal hemipelagic sedimentation took place (Cau) or shallow carbonate deposits (Puentenansa) were recorded (Fig. 13). This distribution could correspond to a model of deep stagnant oxygen-depleted waters (i.e. Jenkyns, 1980; Pedersen and Calvert, 1990). During the C3 interval, an episode of environmental change affected the northern, more restricted NCB, leading to the drowning of the Puentenansa carbonate platform and deposition of organic-rich facies (Najarro et al., 2011a), under oxygen-depleted, probably stratified, waters. Contemporaneously in the SIPM environments, a short episode of organic-rich sediment deposition took place in the shallower Cau sector, whereas in shallow carbonate platforms of the SIPM a drowning event took place (i.e. Castro et al., 2008; Castro et al., 2012). An interesting question is the temporal relationship between platform drowning and the onset of the OAE1a (i.e. Weissert et al., 1998). The data from Puentenansa indicate that the platform drowning predates the C4 segment, which is in agreement with previous



**Fig. 12.** Pr/ $C_{17}$  vs. Ph/ $C_{18}$  plot. The reference lines are from Didyck et al., 1978, but should not be interpreted against the data from less thermally mature sections (CAU and XF) as they have been empirically derived from oils and source rocks.



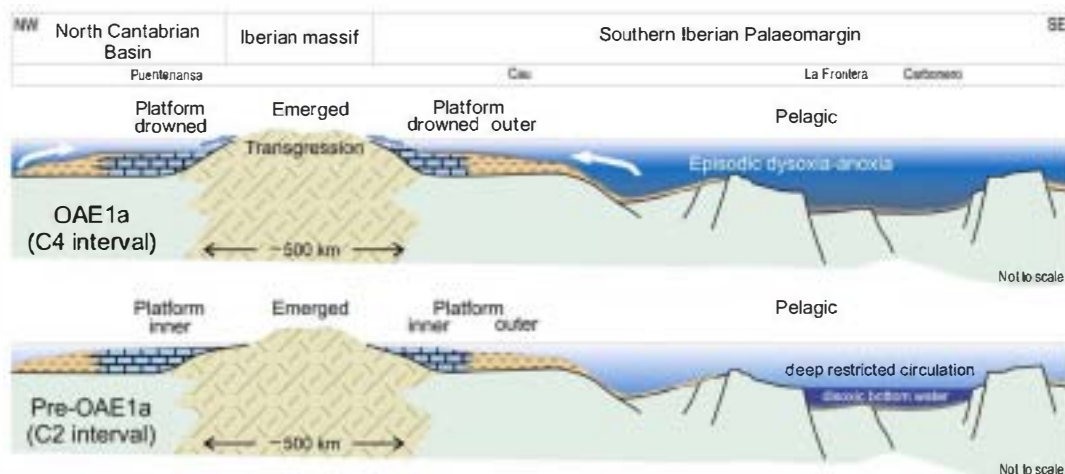


Fig. 13. Sedimentary model. See text for explanation.

studies from the Betic Cordillera (Castro et al., 2012), and also from the Subalpine Chains (Huck et al., 2011).

During the C4–C5 intervals, which corresponds to the global widespread deposition of organic matter of the OAE1a (i.e. Menegatti et al., 1998; Dumitrescu and Brassell, 2006) organic-rich sedimentation characterized all four sections. This persisted until the C8 interval, except at Cau, where organic rich sedimentation ended during C7 interval. This distribution could reflect an expansion of the OMZ, reaching shallower areas on the margins, coincident with a major transgression, which could favor increased continental runoff and an increase in nutrient inputs (i.e. Burla et al., 2008; Castro et al., 2008; Blattler et al., 2011; Najarro et al., 2011a). Similar models of expansion of the OMZ have been proposed for the OAE2 (i.e. Pancost et al., 2004; Li et al., 2006).

The main factors leading to deposition of organic-rich marine sediments are anoxia and productivity (e.g. Pedersen and Calvert, 1990). Marine organic-rich sediments related to OAEs have been extensively interpreted as being deposited under anoxic or dysoxic conditions (e.g. Jenkyns, 1980). In the Iberian sections studied here, evidence for anoxia has been found, both from sedimentology (high TOC contents, barite concretions, laminations, lack of bioturbation, planktonic associations...), and biomarkers (presence of gammacerane, distribution of homohopanes, Pr/Ph ratios, isoprenoids/n-alkane ratios). However, the signature for anoxia is neither unambiguous nor widespread. Differences among sections are interesting: Carbonero section shows a stronger sedimentologic signal of anoxia (i.e. highest TOC contents), whereas the biomarker proxies of these conditions are weak. Although deeper studies are needed about redox proxies for the Carbonero section, a high productivity environment should be also considered; this is consistent with the presence of Ba, which has been proposed as a powerful proxy for productivity in marine environments (Paytan and Griffith, 2007), although more detailed investigation on this element are necessary to avoid misinterpretations. The other SIPM sections (La Frontera and Cau) show both sedimentary and biomarker evidence of anoxia, as stated before. On the other hand, the NCB section (Puentenansa) shows stronger biomarker evidence for anoxia (presence of gammacerane, homohopane distributions, Pr/Ph ratios), which can be related to the palaeogeographic setting in the Bay of Biscay, probably with a more restricted communication of the waters to the open ocean.

Finally, although the sections studied represent different sedimentary environments (platform and pelagic), there are no systematic biomarker differences related to this, for example, there is no evidence for greater terrigenous OM inputs to the platform settings.

Instead, the main differences in biomarker distributions are associated with thermal maturity and vertical variability.

## 6. Conclusions

Four sections recording the Early Aptian OAE1a in Spain have been characterized by biostratigraphy, C-isotope stratigraphy, elemental geochemistry, biomarker distribution, and sedimentology; the sections represent different palaeogeographic (North Cantabrian Basin and Southern Iberian Palaeomargin) and geotectonic settings (pelagic and shallow marine). The C-isotope curves allowed the recognition of the eight segments proposed by Menegatti et al. (1998) and subsequently identified in sections from around the world, and the integration of biostratigraphic and C-stratigraphic data was used to develop an integrated stratigraphic framework, allowing correlation of the Iberian OAE1a. Biomarker distributions revealed significant variations in thermal maturity, which was very low in the Cau and La Frontera sections, moderate in the Puentenansa samples, but relatively high for the Carbonero section, which is within the oil window. The strong lateral variations in maturity over short distances were interpreted as probably due to volcanic activity or burial history. Overall, all of the OM present in the studied samples is interpreted to derive from significant terrestrial inputs as well as marine and bacterial sources, and dinosterane was particularly abundant in several samples. Environmental proxies indicative of anoxia–dysoxia occur in all the sections, but evidence for strong and persistent water column anoxia is equivocal. Crucially, however, the correlation of the sections reveals that deposition of organic-rich facies started earlier in pelagic settings and later in the platform settings, which can be related to an expansion of the oxygen minimum zone from deep marine waters to shallower marine environments during the development of the OAE1a.

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